AMENDMENTS TO THE CLAIMS:

The listing of claims shown below will replace all prior versions, and listings, of claims in the Application:

Claims 1-48 (Cancelled)

Claim 49. (Currently Amended) A method of managing the pharmaceutical care of a patient using one or more software-accessible databases comprising the steps of:

providing a clinical database on a plurality of drugs, each drug in the database being associated with a multi-character therapeutic cross reference code, wherein a first set of characters represent a class of drugs, a second set of characters represents a subclass of drugs, and third set of characters represent a specific drug;

updating a patient database with a drug therapy regimen for the patient, the drug therapy regimen comprising an identification of each drug prescribed to the patient, a frequency per day for each drug, and a daily dosage for each drug;

updating the patient database with patient data, the patient data comprising any disease states and allergies for the patient;

querying [[a]] the clinical database with the drug therapy regimen and patient data, wherein the querying step further comprises identifying: (a) allergies the patient has for any of the prescribed drugs; (b) drug-drug interactions for any of the prescribed drugs; (c) dosage irregularities; (d) drug-disease contraindications; (e) therapeutic duplications; (f) drug(s) in the drug therapy regimen without a medical indication; (g) adverse drug reactions; and (h) untreated disease states wherein the identification is based at least in

part on a comparison of the multi-character therapeutic cross reference code with the patient database records; and

generating a report based on the querying step.

- Claim 50. (Previously Presented) The method according to claim 49, wherein the querying step identifies the following additional information for each patient:
- (i) information regarding use or efficacy of any of the prescribed drugs; and
 - (j) information regarding patient compliance.
- Claim 51. (Previously Presented) The method according to claim 50, wherein the querying step identifies the following additional information for each patient:
- (k) information regarding an assessment of the educational needs of the patient; and
 - (I) information regarding the financial circumstances of the patient.
- Claim 52. (Previously Presented) The method according to claim 49, wherein the drug therapy regimen for the patient comprises a plurality of drugs prescribed by more than one physician.
- Claim 53. (Previously Presented) The method according to claim 49, further comprising the step of modifying the drug therapy regimen based on the report.

Claim 54. (New) A method of managing the pharmaceutical care of a patient using one or more software-accessible databases comprising the steps of:

providing a clinical database on a plurality of drugs, each drug in the database being associated with a multi-character therapeutic cross reference code, wherein a first set of characters represent a class of drugs, a second set of characters represents a subclass of drugs, and third set of characters represent a specific drug;

updating a patient database with a drug therapy regimen for the patient, the drug therapy regimen;

updating the patient database with patient data, the patient data comprising any disease states and allergies for the patient;

identifying: (a) allergies the patient has for any of the prescribed drugs; (b) drugdrug interactions for any of the prescribed drugs; (c) dosage irregularities; (d) drugdisease contraindications; (e) therapeutic duplications; (f) drug(s) in the drug therapy regimen without a medical indication; (g) adverse drug reactions; and (h) untreated disease states based at least in part on a comparison of the multi-character therapeutic cross reference code with the patient database records.

Claim 55. (New) The method of claim 54, wherein the multi-character therapeutic cross reference code comprises an eight character code with the first two characters represent a class of drugs, the next four characters represent a subclass of drugs, and the next two characters represent a specific drug.

Claim 56. (New) The method of claim 54, wherein the multi-character therapeutic

cross reference code is associated with drug indications and contra-indications via ICD-9 codes.

- Claim 57. (New) The method of claim 54, further comprising the step of generating a report.
- Claim 58. (New) The method of claim 54, wherein the patient database is updated with drug therapy regimen data and a compliance percentage is generated.
- Claim 59. (New) The method of claim 54, wherein the drug therapy regimen data is automatically imported from a pharmacy dispensing system.
- Claim 60. (New) A method for identifying one or more drugs causing an identified adverse reaction using one or more software-accessible databases, comprising the steps of:

providing a clinical database on a plurality of drugs, each drug in the database being associated with a multi-character therapeutic cross reference code, wherein a first set of characters represent a class of drugs, a second set of characters represents a subclass of drugs, and third set of characters represent a specific drug, the clinical database further including adverse reaction information associated with each drug;

querying the clinical database with a given adverse reaction;

identifying all drugs in a class or subclass of drugs having the given adverse reaction based at least in part on the multi-character therapeutic cross reference code.

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Claim 61. (New) The method of claim 60, wherein the identifying step highlights a particular drug in a patient's current drug regimen in addition to listing other drugs in the class or subclass with the same adverse reaction.